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Amendments to Claims:

This listing of claims will replace all prior versions and listings in the application:

Listing of Claims:

- 1-98. (Canceled)
- 99. (Previously Presented) The pharmaceutical composition according to claim 116, wherein the dendritic cells express an amount of the modified antigen to provide between about 1 to 100 micrograms of the modified antigen in said pharmaceutical composition.
 - 100. (Canceled)
- 101. (Currently Amended) An *in vitro* composition comprising mature dendritic cells expressing modified antigen and derived from an *in vitro* culture of an enriched and expanded population of proliferating dendritic cell precursors by a method comprising:

providing a tissue source comprising dendritic cell precursors;

treating the tissue source comprising dendritic cell precursors to increase the proportion of dendritic cell precursors;

treating the tissue source to obtain a population of cells suitable for culture in vitro;

culturing the tissue source on a substrate in a culture medium comprising GM-CSF to obtain cell aggregates comprising proliferating dendritic cell precursors nonadherent cells and cell clusters;

subculturing the cell aggregates at least one time to enrich the proportion of the nonadherent cells and cell clusters to produce cell aggregates comprising proliferating dendritic cell precursors;

serially subculturing the cell aggregates one or more times to enrich the proportion of dendritic cell precursors; and

continuing to culture the dendritic cell precursors for a period of time to allow them to mature into mature dendritic cells;

wherein the dendritic cells are cultured culturing the dendritic cells in vitro in the presence of an antigen for a time sufficient to allow the antigen to bind to

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the dendritic cells and, wherein the dendritic cells process the antigen to produce a modified antigen which is expressed by the dendritic cells.

102-103. (Canceled)

- 104. (Previously Presented) The composition according to claim 101, wherein the tissue source is blood.
- 105. (Previously Presented) The composition according to claim 101, wherein the tissue source is bone marrow.
- 106. (Previously Presented) The composition according to claim 101, wherein GM-CSF is present in the culture medium at a concentration of about 1-1000 U/ml.
- 107. (Previously Presented) The composition according to claim 104, wherein the concentration of GM-CSF in the culture medium is about 30-100 U/ml.
- 108. (Previously Presented) The composition according to claim 105, wherein the concentration of GM-CSF in the culture medium is about 500-1000 U/ml.
- 109. (Previously Presented) The composition according to claim 101, wherein the cell aggregates are blood derived and are subcultured from about one to five times.
- 110. (Previously Presented) The composition according to claim 101, wherein the cell aggregates are subcultured one to five times.
- 111. (Previously Presented) The composition according to claim 101, wherein the culture medium is selected from the group consisting of RPMI 1640, DMEM and α -MEM, and wherein the culture medium is supplemented with scrum.
- 112. (Previously Presented) The composition according to claim 104, wherein the tissue source is treated to remove red blood cells.
- 113. (Previously Presented) The composition according to claim 105, wherein the tissue source is treated to remove B cells and granulocytes.

114-115. (Canceled)

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116. (Previously Presented) A pharmaceutical composition comprising a therapeutically effective amount of the composition according to claim 101.

117-119. (Canceled)

- expressing a modified antigen and derived from an *in vitro* culture of a population of enriched and expanded proliferating dendritic cell precursor cells, wherein said dendritic cells are contacted *in vitro* with antigen in the presence of GM CSF for a sufficient time to allow the antigen to bind to the dendritic cells; by a method comprising culturing dendritic cell precursor cells in a culture medium comprising GM-CSF at a concentration sufficient to promote the survival and proliferation of dendritic cell precursors; serially subculturing the proliferating dendritic cell precursors at intervals which provide for the continued proliferation of said dendritic cell precursors; and exposing the cells to antigen *in vitro*, wherein the dendritic cells process the antigen to produce a modified antigen which is expressed by the dendritic cells.
 - 121-141. (Cancelled)
- 142. (Previously Presented) The composition according to claim 101, wherein the dendritic cell precursors are human.
- 143. (Currently Amended) The composition of dendritic cell procursors according to claim 142, wherein the dendritic cell precursors are obtained from blood.
- 144. (Currently Amended) The composition of dendritic cell precursors according to claim 142, wherein the dendritic cell precursors are obtained from bone marrow.
- 145. (New) An *in vitro* composition comprising antigen-activated dendritic cells expressing modified antigens and derived from an *in vitro* culture of proliferating dendritic cell precursors by a method comprising:
 - a) providing a tissue source comprising dendritic cell precursors;
 - b) treating the tissue source to obtain a population of cells suitable for culture in vitro;
 - c) culturing the tissue source on a substrate in a culture medium comprising GM-CSF to obtain nonadherent cells and cell clusters;

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- d) subculturing the nonadherent cells and cell clusters to produce cell aggregates comprising proliferating dendritic cell precursors;
- e) serially subculturing the cell aggregates one or more time to enrich the proportion of dendritic cell precursors;
- f) continuing to culture the dendritic cell precursors for a period of time sufficient to allow them to mature into mature dendritic cells; and
- g) pulsing the dendritic cells with an antigen, wherein the dendritic cells process the antigen to produce a modified antigen which is expressed by the dendritic cells.